

AD-A276 668

94-07073

REPORT DOCUMENTATION PAGE

Form Approved
OMB No 0704-0188

2

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302 and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503.

1. AGENCY USE ONLY (Leave blank)		2. REPORT DATE 25-2-94	3. REPORT TYPE AND DATES COVERED Technical Report
4. TITLE AND SUBTITLE Use of Organoindium Hydrides for the Preparation of Organoindium Phosphides. Synthesis and Molecular Structure of $[(Me_3CCH_2)_2InP(t-Bu)_2]_2$			5. FUNDING NUMBERS Grant: N00014-90-J-1530 R&T Code: 4135002
6. AUTHOR(S) O.T. Beachley, Jr., Sun-Hua L. Chao, Melvyn Rowen Churchill and Charles H. Lake			8. PERFORMING ORGANIZATION REPORT NUMBER Technical Report No. 37
7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) Department of Chemistry State University of New York at Buffalo Buffalo, NY 14214-3094			10. SPONSORING/MONITORING AGENCY REPORT NUMBER n/a
9. SPONSORING/MONITORING AGENCY NAME(S) AND ADDRESS(ES) Office of Naval Research 800 N. Quincy Street Arlington, VA 22217-5000			11. SUPPLEMENTARY NOTES Accepted for publication - Organometallics
12a. DISTRIBUTION/AVAILABILITY STATEMENT This document has been approved for public release and sale; its distribution is unlimited. Reproduction in whole or in part is permitted for any purpose of the United States government.			12b. DISTRIBUTION CODE n/a
13. ABSTRACT (Maximum 200 words) The indium phosphide $[(Me_3CCH_2)_2InP(t-Bu)_2]_2$ has been prepared from $K[In(CH_2CMe_3)_3H]$ and $ClP(t-Bu)_2$ in pentane. When $In(CH_2CMe_3)_3$ and $HP(t-Bu)_2$ were present in a 1:1 mol ratio, heating to 105-115 °C for 5 days was required, whereas when $In(CH_2CMe_3)_3$ and $HP(t-Bu)_2$ were in a 5:1 mol ratio in pentane solution, large, colorless crystal of the desired indium product formed in 6 days at room temperature. Excess phosphine, $In(CH_2CMe_3)_3$ and $HP(t-Bu)_2$ in a 1:5 mol ratio in pentane, significantly retarded the rate of formation of $[(Me_3CCH_2)_2InP(t-Bu)_2]_2$. Thermal decomposition of $[(Me_3CCH_2)_2InP(t-Bu)_2]_2$ to form InP occurred at 245 °C in 1 h. The compound $[(Me_3CCH_2)_2InP(t-Bu)_2]_2$ crystallizes in the centrosymmetric orthorhombic space group $Pbcn$ (No. 60) with $a = 11.742(3)\text{\AA}$, $b = 20.194(6)\text{\AA}$, $c = 17.909(4)\text{\AA}$, $V =$ (continued on next page)			
14. SUBJECT TERMS Indium-phosphorus compounds, indium-hydrides, single-source precursor, X-ray structural study			15. NUMBER OF PAGES 26
			16. PRICE CODE n/a
17. SECURITY CLASSIFICATION OF REPORT Unclassified	18. SECURITY CLASSIFICATION OF THIS PAGE Unclassified	19. SECURITY CLASSIFICATION OF ABSTRACT Unclassified	20. LIMITATION OF ABSTRACT UL

NSN 7540-01-280-5500

Standard Form 298 (Rev 2-89)
Prescribed by ANSI Std Z39-18
298-102

94 3 03 013

4246(2) \AA^3 and $Z = 4$. The structure was solved and refined to $R = 6.68\%$ and $wR = 6.24\%$ for all 4920 independent reflections and $R = 2.76\%$ and $wR = 3.47\%$ for those 2525 reflections with $|F_o| > 6.0\sigma(|F_o|)$. The molecule lies on a two-fold axis which passes through the two indium atoms and requires that the In_2P_2 core be strictly planar.

Accession For	
NTIS	CRA&I <input checked="" type="checkbox"/>
DTIC	TAB <input checked="" type="checkbox"/>
U. announced <input type="checkbox"/>	
Justification _____	
By _____	
Distribution / _____	
Availability Codes	
Dist	Avail and/or Special
A-1	

DTIC QUALITY INSPECTED 2

OFFICE OF NAVAL RESEARCH

Contract N-00014-90-J-1530

R&T Code 4135002

TECHNICAL REPORT NO. 37

Use of Organoindium Hydrides for the Preparation of
Organoindium Phosphides. Synthesis and Molecular
Structure of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$

by

O. T. Beachley, Jr., Sun-Hua L. Chao, Melvyn Rowen Churchill
and Charles H. Lake

Prepared for Publication

in

Organometallics

State University of New York at Buffalo
Department of Chemistry
Buffalo, New York 14214

25 February 1994

Reproduction in whole or in part is permitted for
any purpose of the United States Government

*This document has been approved for public release
and sale; its distribution is unlimited

Contribution from the Department of Chemistry
State University of New York at Buffalo, Buffalo, NY 14214

Use of Organoindium Hydrides for the Preparation
Of Organoindium Phosphides. Synthesis and
Molecular Structure of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$

by

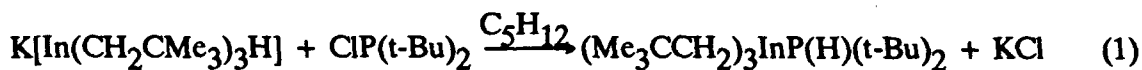
O. T. Beachley, Jr.*, Sun-Hua L. Chao, Melvyn Rowen
Churchill* and Charles H. Lake

Summary: The indium phosphide $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ has been prepared from $\text{K}[\text{In}(\text{CH}_2\text{CMe}_3)_3\text{H}]$ and $\text{ClP}(\text{t-Bu})_2$ in pentane. When $\text{In}(\text{CH}_2\text{CMe}_3)_3$ and $\text{HP}(\text{t-Bu})_2$ were present in a 1:1 mol ratio, heating to 105-115 °C for 5 days was required, whereas when $\text{In}(\text{CH}_2\text{CMe}_3)_3$ and $\text{HP}(\text{t-Bu})_2$ were in a 5:1 mol ratio in pentane solution, large, colorless crystal of the desired indium product formed in 6 days at room temperature. Excess phosphine, $\text{In}(\text{CH}_2\text{CMe}_3)_3$ and $\text{HP}(\text{t-Bu})_2$ in a 1:5 mol ratio in pentane, significantly retarded the rate of formation of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$. Thermal decomposition of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ to form InP occurred at 245 °C in 1 h. The compound $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ crystallizes in the centrosymmetric orthorhombic space group Pbcn (No. 60) with $a = 11.742(3)\text{\AA}$, $b = 20.194(6)\text{\AA}$, $c = 17.909(4)\text{\AA}$, $V = 4246(2)\text{\AA}^3$

and $Z = 4$. The structure was solved and refined to $R = 6.68\%$ and $wR = 6.24\%$ for all 4920 independent reflections and $R = 2.76\%$ and $wR = 3.47\%$ for those 2525 reflections with $|F_o| > 6.0\sigma(|F_o|)$. The molecule lies on a two-fold axis which passes through the two indium atoms and requires that the In_2P_2 core be strictly planar.

Compounds of the type R_2MER_2' (M = group 13 element, E = group 15 element) are OMCVD precursors for the preparation of group 13-15 materials such as GaAs^{1,3-5} and InP². The original and simplest synthetic route to R_2InPR_2' is probably a hydrocarbon elimination reaction⁶⁻¹³ between InR_3 and HPR_2' . Metathesis reactions^{9,10,14} are also useful for the preparation of R_2InPR_2' but require preparation of InR_2X (X = Cl, Br, I) and $M'ER_2'$ (M = Li, Na, K). Similarly, the Me_3SiCl elimination reaction^{15,16} requires the syntheses of InR_2X and $(Me_3Si)PR_2'$. A goal of our research has been to develop a scheme for the synthesis of R_2InPR_2' which would take advantage of the simplicity of the elimination reaction but which would not require the prior synthesis and handling of HPR_2' . Ideally, the secondary phosphine should be formed and then consumed by reaction with the organoindium compound. These concepts have been applied successfully to the synthesis of $[(Me_3CCH_2)_2InP(t-Bu)_2]_2$ from $K[In(CH_2CMe_3)_3H]$ ¹⁷ and $ClP(t-Bu)_2$ in pentane. The new indium phosphide derivative has been prepared in high yield and has been fully characterized but it was too insoluble in benzene for a cryoscopic molecular weight study.

The reaction of $K[In(CH_2CMe_3)_3H]$ and $ClP(t-Bu)_2$ leads to the formation of $In(CH_2CMe_3)_3$ and $HP(t-Bu)_2$ (Equation 1) in high, if not quantitative, yield. These

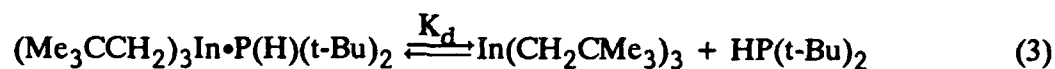


reaction products subsequently undergo the elimination reaction under appropriate conditions to form $[(Me_3CCH_2)_2InP(t-Bu)_2]_2$ and neopentane (Equation 2). When

$\text{In}(\text{CH}_2\text{CMe}_3)_3$ and $\text{HP}(\text{t-Bu})_2$ were present in a 1:1 mol ratio, heating to 105-115 °C for 5 days was necessary to effect the formation of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ in high (~90%) yield. However, when $\text{In}(\text{CH}_2\text{CMe}_3)_3$ and $\text{HP}(\text{t-Bu})_2$ were in a 5:1 mol ratio in pentane solution, large, colorless crystals of the desired indium product formed in 6 days at room temperature. When $\text{In}(\text{CH}_2\text{CMe}_3)_3$ and $\text{HP}(\text{t-Bu})_2$ were present in a 1:5 mol ratio in pentane solution, no $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ was formed, even after a month. Apparently, excess $\text{In}(\text{CH}_2\text{CMe}_3)_3$ enhances the rate of the elimination reaction. The simple adduct $(\text{Me}_3\text{CCH}_2)_3\text{InP}(\text{H})(\text{t-Bu})_2$ probably is not involved in the actual elimination reaction at room temperature. If the simple 1:1 adduct had kinetic significance for the elimination reaction, excess $\text{In}(\text{CH}_2\text{CMe}_3)_3$ as well as excess $\text{HP}(\text{t-Bu})_2$ should have increased the rate of elimination similarly. It is also noteworthy that $\text{K}[\text{In}(\text{CH}_2\text{CMe}_3)_3\text{H}]$ does not deprotonate preformed $\text{HP}(\text{t-Bu})_2$ to form $\text{KP}(\text{t-Bu})_2\text{H}_2$ and $\text{In}(\text{CH}_2\text{CMe}_3)_3$.

The elimination reactions between stoichiometric quantities of $\text{HP}(\text{t-Bu})_2$ and InMe_3 ^{11,13} and InEt_3 ¹² have been reported previously. The neat reagents InMe_3 and $\text{HP}(\text{t-Bu})_2$ required heating¹¹ to 170 °C for 6 h to effect formation of $[\text{Me}_2\text{InP}(\text{t-Bu})_2]_2$ whereas the preparation of $[\text{Et}_2\text{InP}(\text{t-Bu})_2]_2$ utilized a refluxing hexane solution (~70 °C).¹²

The nature of the simple adduct $(\text{Me}_3\text{CCH}_2)_3\text{In}\cdot\text{P}(\text{H})(\text{t-Bu})_2$ has been investigated by freezing point depression studies in benzene and by ¹H and ³¹P NMR spectroscopy. The equilibrium constant for the dissociation of the adduct in benzene has been calculated from cryoscopic data at -5 °C to be 0.011 ± 0.002 (K_d), which is



comparable to the K_d of 9.1×10^{-3} for the dissociation of $\text{HMe}_2\text{Al}\cdot\text{N}(\text{Me})(\text{Ph})(\text{H})$ ¹⁸ at -63 °C. The K_d of the adduct $\text{HMe}_2\text{Al}\cdot\text{P}(\text{Me})(\text{Ph})(\text{H})$ ¹⁹ is 0.38 at 22 °C. The NMR

spectroscopic data for $(\text{Me}_3\text{CCH}_2)_3\text{In}\cdot\text{P}(\text{H})(\text{t-Bu})_2$ indicate rapid exchange between the components of the equilibrium. Consequently, the ^1H and ^{31}P NMR chemical shifts and coupling constant data depend upon the phosphorus to indium ratio. Extrapolation of these data to $\text{HP}(\text{t-Bu})_2/\text{In}(\text{CH}_2\text{CMe}_3)_3$ of zero provided nominal data for pure adduct as $^{31}\text{P}(\delta)$ 14.70 ppm. (dm, $^1J = 269$ Hz, $^3J = 12.9$ Hz) and $^1\text{H}(\delta)$ 3.24 ppm (d, PH, $^1J = 269$ Hz). It is also of interest that the adduct $(\text{Me}_3\text{CCH}_2)_3\text{In}\cdot\text{P}(\text{H})(\text{t-Bu})_2$ reacts with KH to form $\text{K}[\text{In}(\text{CH}_2\text{CMe}_3)_3\text{H}]$ and free $\text{HP}(\text{t-Bu})_2$. Thus, $\text{HP}(\text{t-Bu})_2$ can be readily isolated from the reaction of $\text{K}[\text{In}(\text{CH}_2\text{CMe}_3)_3\text{H}]$ with $\text{ClP}(\text{t-Bu})_2$, if excess KH is present.

A single-crystal X-ray structural study of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ shows that this compound consists of discrete molecular units of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ separated by normal van der Waals distances. The labelling of atoms is provided in Figure 1. Distances and angles are collected in Table I. The molecule has precise C_2 symmetry, with atoms In(1) and In(2) lying on a crystallographic two-fold axis. This requires that the In_2P_2 core be strictly planar. Indium-phosphorus distances are $\text{In}(1)\text{-P}(1) = \text{In}(1)\text{-P}(1\text{A}) = 2.712(1)\text{\AA}$ and $\text{In}(2)\text{-P}(1) = \text{In}(2)\text{-P}(2\text{A}) = 2.690(1)\text{\AA}$ (average = 2.701\AA). Internal angles at indium are $\text{P}(1)\text{-In}(1)\text{-P}(1\text{A}) = 83.0(1)^\circ$ and $\text{P}(1)\text{-In}(2)\text{-P}(1\text{A}) = 83.8(1)^\circ$ [average $\text{P-In-P} = 83.4^\circ$]; internal angles at phosphorus are symmetry equivalent, with $\text{In}(1)\text{-P}(1)\text{-In}(2) = \text{In}(1)\text{-P}(1\text{A})\text{-In}(2) = 96.6(1)^\circ$. Cross-ring distances are clearly non-bonding, with $\text{In}(1)\cdots\text{In}(2) = 4.033\text{\AA}$ and $\text{P}(1)\cdots\text{P}(1\text{A}) = 3.515\text{\AA}$.

Two neopentyl groups are associated with each indium atom, with indium-carbon distances of $\text{In}(1)\text{-C}(31) = \text{In}(1)\text{-C}(31\text{A}) = 2.206(5)\text{\AA}$ and $\text{In}(2)\text{-C}(41) = \text{In}(2)\text{-C}(41\text{A}) = 2.219(4)\text{\AA}$ (average $\text{In-C} = 2.213\text{\AA}$). Interligand angles are $\text{C}(31)\text{-In}(1)\text{-C}(31\text{A}) = 117.7(2)^\circ$ and $\text{C}(41)\text{-In}(2)\text{-C}(41\text{A}) = 112.7(2)^\circ$. The $\text{P-In-C}(\text{neopentyl})$ angles are not equivalent.

Thus $\text{P}(1)\text{-In}(1)\text{-C}(31) = 116.6(1)^\circ$ as compared to $\text{P}(1\text{A})\text{-In}(1)\text{-C}(31) = 109.1(1)^\circ$ and $\text{P}(1\text{A})\text{-In}(2)\text{-C}(41) = 119.2(1)^\circ$ as compared to $\text{P}(1)\text{-In}(2)\text{-C}(41) = 109.7(1)^\circ$. The C-In-C planes are therefore not precisely perpendicular to the In_2P_2 core. Thus, the $\text{C}(31)\text{-In}(1)\text{-C}(31\text{A})$ plane is oriented at 83.9° to the In_2P_2 core while the $\text{C}(41)\text{-In}(2)\text{-C}(41\text{A})$ plane is oriented at 97.8° to the In_2P_2 core. We note here that the neopentyl groups attached to $\text{In}(2)$ are well behaved with normal vibration ellipsoids (see Figure 1) and normal carbon-carbon bond lengths ($\text{C}(41)\text{-C}(42) = 1.552(6)\text{\AA}$ and $\text{C}(42)\text{-CH}_3$ distances of $1.538(6)$, $1.529(6)$ and $1.530(6)\text{\AA}$). In contrast to this, the neopentyl groups attached to $\text{In}(1)$ are associated with substantial librational motion about the $\text{C}(31)\text{-C}(32)$ axis, which results in artificial librational shortening of the $\text{C}(32)\text{-CH}_3$ bonds (i.e., $\text{C}(31)\text{-C}(32) = 1.539(6)\text{\AA}$ but $\text{C}(32)\text{-C}(33) = 1.455(8)\text{\AA}$, $\text{C}(32)\text{-C}(34) = 1.449(8)\text{\AA}$ and $\text{C}(32)\text{-C}(35) = 1.457(7)\text{\AA}$).

The $\text{P}(\text{t-Bu})_2$ ligands with phosphorus - carbon distances of $\text{P}(1)\text{-C}(11) = 1.895(4)\text{\AA}$ and $\text{P}(1)\text{-C}(21) = 1.891(4)\text{\AA}$ (average $\text{P-C} = 1.893\text{\AA}$) also are not strictly perpendicular to the In_2P_2 plane. The $\text{C}(11)\text{-P}(1)\text{-C}(21)$ plane makes an angle of 88.6° with the In_2P_2 plane. The In-P-C angles are inequivalent, with $\text{In}(1)\text{-P}(1)\text{-C}(11) = 116.7(1)^\circ$ as compared to $\text{In}(1)\text{-P}(1)\text{-C}(21) = 109.6(1)^\circ$ and $\text{In}(2)\text{-P}(1)\text{-C}(21) = 112.6(1)^\circ$ as compared to $\text{In}(2)\text{-P}(1)\text{-C}(11) = 110.6(1)^\circ$.

The structure of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ can be compared with the closely related structures of $[\text{Me}_2\text{InP}(\text{t-Bu})_2]_2$ ¹¹ and $[\text{Et}_2\text{InP}(\text{t-Bu})_2]_2$.¹² Important distances and angles are compared in Table II. The indium-phosphorus and indium-carbon distances in the neopentyl derivative are significantly longer than those in the methyl and ethyl compounds. It is also noteworthy that the indium atoms are further apart in the neopentyl derivative because the P-In-P angles are the smallest of the three compounds. This decreased P-In-P

angle might be the result of the larger C-In-C angle needed to keep the more sterically demanding neopentyl ligands apart. The In-P-In angles are also largest for the neopentyl derivative.

The complete characterization of an organoindium phosphide requires the determination of the degree of association of the compound in the solid, solution and gas phases, if possible. The X-ray structural study demonstrated the existence of the dimer $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ as the solid. The low solubility of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ in benzene solution prevented cryoscopic molecular weight studies. Thus, none of the three known t-butylphosphide derivatives are sufficiently soluble in benzene for molecular weight studies even though all are simple dimers in the solid state.^{11,12} The methyl derivative¹¹ $\text{Me}_2\text{InP}(\text{t-Bu})_2$ was even too insoluble in benzene for NMR studies. Even though molecular weight studies were impossible, the ^1H NMR spectrum of $(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2$ is consistent with the existence of dimers in benzene solution. The resonances for the t-butyl group protons are two overlapping doublets which appear like a triplet. The t-butyl protons are thus coupled to two phosphorus atoms in the four-membered ring. The mass spectrum of $(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2$ suggests the existence of monomers in the gas phase. In contrast, the mass spectrum of $\text{Me}_2\text{InP}(\text{t-Bu})_2$ demonstrated the presence of dimers.¹¹

The thermolysis of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ at 245 °C for 1 h in a sealed tube leads to the formation of InP with relatively low contamination by carbon and hydrogen (0.41% C and 0.04% H). The identity of the black solid remaining in the tube as InP was verified by X-ray powder diffraction data and an XPS spectrum. Good quality InP has also been grown from $[\text{Me}_2\text{InP}(\text{t-Bu})_2]_2$ at 350-480 °C under OMCVD conditions.² Preliminary data indicated that $[\text{Me}_2\text{InP}(\text{t-Bu})_2]_2$ decomposed at 240 °C whereas $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$

decomposed at a lower temperature, 200-202 °C. The lower temperature required for $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ might be due to its more facile dissociation to monomers. The more bulky neopentyl groups provide longer and supposedly weaker indium-phosphorus ring bonds which could make the monomer more accessible in the gas phase. The monomer with vacant orbitals should in turn decompose more readily than the dimer. The mass spectra of the two compounds suggest that the monomer of the neopentyl derivative is readily formed in the gas phase. The compound $[\text{Me}_2\text{InP}(\text{t-Bu})_2]_2$ ¹¹ exhibited masses of species associated with dimers and monomers, whereas no species related to the dimer of the neopentyl derivative were observed.

Experimental

All compounds described in this investigation are extremely sensitive to oxygen and moisture and were manipulated in a standard vacuum line or under a purified argon atmosphere in a Vacuum/Atmospheres drybox. The starting compounds, $\text{In}(\text{CH}_2\text{CMe}_3)_3$ ²⁰ and $\text{K}[\text{In}(\text{CH}_2\text{CMe}_3)_3\text{H}]$ ¹⁷ were prepared and purified by literature methods. The chlorophosphine $\text{ClP}(\text{t-Bu})_2$ was purchased from Strem Chemicals, Inc. and distilled twice before use. The reagent KH was obtained from Aldrich Chemical Co. and was washed with pentane to remove oil prior to use. Solvents were dried by conventional procedures. Elemental analyses for $(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2$ and InP were performed by Schwarzkopf Microanalytical Laboratory, Woodside, NY and by E+R Microanalytical Laboratory, Inc., Corona, NY, respectively. Infrared spectra of Nujol mulls between CsI plates were recorded by means of a Perkin-Elmer 683 spectrometer. The ^1H NMR spectra were recorded at 400 MHz by means of a Varian VXR-400 S spectrometer, or at 300 MHz with a Varian Gemini-300 spectrometer. Proton chemical shifts are reported in δ units (ppm) and

are referenced to SiMe_4 at 0.00 ppm and C_6H_6 at δ 7.15 ppm. The ^{31}P NMR spectra were recorded at 161.9 MHz by means of a Varian VXR-400 S spectrometer. The ^{31}P spectra are referenced to 85% H_3PO_4 at δ 0.00 ppm. All samples for NMR spectra were contained in sealed NMR tubes. Melting points were observed in sealed capillaries filled with purified argon and are uncorrected. Cryoscopic studies of $(\text{Me}_3\text{CCH}_2)_3\text{In}\cdot\text{P}(\text{H})(\text{t-Bu})_2$ in benzene were obtained by using an instrument similar to that described by Shriver and Drezdson.²¹ Mass spectrometry was conducted by using a VG Analytical 70-SE spectrometer. Samples were sealed in microcapillaries filled with purified argon which were broken open just before insertion into the chamber of the spectrometer. Species were ionized by electron impact at 70 eV with the source temperature at 200 °C. Only peaks with m/z greater than 100 and with I_{rel} higher than 10 are reported. Masses of species containing In refer to the ^{115}In isotope. The XPS analysis of InP was performed on a Perkin-Elmer Physical Electronics (PHI) Model 5100 ESCA spectrometer with a $\text{Mg K}\alpha_{1,2}$ X-ray source (1253.6 eV) operated at 300 W, 15 kV and 20 mA, and was recorded at a take-off angle of 45° with a 180° hemispherical detector. The spectrometer was calibrated at a base pressure of 2×10^{-9} torr and at an operating pressure of 2×10^{-8} torr with the $\text{Ag } 3d_{5/2}$ peak set at 367.9 eV with FWHM (full width at half maximum) of 1.05 and 900,000 counts per second. The binding energy scale was set by $\text{Cu } 2p_{3/2}$ and $\text{Cu } 3p_{3/2}$ at 932.5 and 75.0 eV, respectively. The identity of the elements was evaluated by low-resolution (89.45 eV) survey spectra. Binding energies (eV) and peak area for quantitative analysis were recorded by high resolution (35.75 eV) spectra. X-ray powder diffraction analysis of InP was recorded on a Siemens D - 500 X-ray diffractometer operated at 40 kV and 30 mA with a graphite

monochromatic Cu $K\alpha$ X-ray source at 1.54 Å. The X-ray powder pattern spectrum was generated as 2θ vs. intensity, and translated into d values.

Synthesis of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$. A solution of $\text{ClP}(\text{t-Bu})_2$ (3.516 g, 19.46 mmol, 15 mL pentane) was slowly added to a solution of $\text{K}[\text{In}(\text{CH}_2\text{CMe}_3)_3\text{H}]$ (7.181 g, 19.49 mmol, 30 mL pentane) at -40 – -50 °C. The mixture was allowed to warm to room temperature slowly, stirred overnight and then filtered. The insoluble KCl (0.963 g, 12.92 mmol, 66% yield based on $\text{ClP}(\text{t-Bu})_2$) was isolated after 8 extractions with pentane. Pentane was then removed from the filtrate at -20 °C by vacuum distillation. The resulting colorless liquid was finally heated at 105 – 115 °C for 5 d. Neopentane (1.239 g, 17.17 mmol, 88.23% based on $\text{ClP}(\text{t-Bu})_2$) was collected by vacuum distillation. The resulting colorless solid was washed 3 times with a small amount of pentane to leave $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ (5.702 g, 7.09 mmol, 73% yield based on $\text{ClP}(\text{t-Bu})_2$) as a colorless crystalline solid. $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$. mp 200 – 202 °C dec. Anal. Calcd for $\text{C}_{18}\text{H}_{40}\text{InP}$: C, 53.74; H, 10.02; P, 7.70. Found: C, 53.75; H, 9.72; P, 7.26. ^1H NMR (C_6D_6): δ 1.39(s, 18 H, InCCCH_3), 1.46 (t, 18 H, PCCH_3 , $J = 6.6$ Hz), 1.54 (s, 4 H, InCH_2). $^{31}\text{P}\{^1\text{H}\}$ NMR (C_6D_6): δ 55.0 (s, PCCH_3). IR (Nujol mull, cm^{-1}) 1360 (vs, sh), 1226 (m), 1207 (w), 1166 (m), 1113 (m, sh), 1012 (m), 927 (vw), 809 (w), 737 (m), 653 (m), 617 (m), 554 (w), 467 (vw), 450 (w). MS m/z (I_{rel}), 402 (13, $(\text{Me}_3\text{CCH}_2)^+\text{InP}(\text{t-Bu})_2^+$), 332 (13, $(\text{Me}_3\text{CCH}_2)(\text{H})\text{InP}(\text{t-Bu})_2^+$), 331 (43, $(\text{Me}_3\text{CCH}_2)\text{InP}(\text{t-Bu})_2^+$), 260 (37, $\text{InP}(\text{t-Bu})_2^+$), 258 (53, $\text{In}(\text{H})(\text{CH}_2\text{CMe}_3)^+$), 257 (97, $\text{In}(\text{CH}_2\text{CMe}_3)_2^+$), 255 (31, $\text{In}(\text{CH}_2\text{CMe}_3)(\text{C}_5\text{H}_9)^+$), 201 (16, $\text{In}(\text{CH}_2\text{CMe}_3)(\text{Me})^+$), 186 (30, $\text{In}(\text{CH}_2\text{CMe}_3)^+$), 146 (61,

InP or $\text{HP}(\text{t-Bu})_2^+$, 145 (14, $\text{P}(\text{t-Bu})_2^+$), 115 (100, In^+ or $\text{P}(\text{C}_3\text{H}_7)(\text{C}_3\text{H}_5)^+$), 113 (34, $\text{P}(\text{C}_3\text{H}_5)_2^+$).

Synthesis of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ with Excess $\text{In}(\text{CH}_2\text{CMe}_3)_3$ at Room

Temperature. The synthesis of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ from $\text{ClP}(\text{t-Bu})_2$ (0.964 g, 5.34 mmol), $\text{K}[\text{In}(\text{CH}_2\text{CMe}_3)_3\text{H}]$ (1.974 g, 5.358 mmol) and excess $\text{In}(\text{CH}_2\text{CMe}_3)_3$ (1.755 g, 5.345 mmol) dissolved in 15 mL pentane was also accomplished at room temperature. After the solutions of $\text{ClP}(\text{t-Bu})_2$ and $\text{K}[\text{In}(\text{CH}_2\text{CMe}_3)_3\text{H}]$ were combined at $-40 \sim -50^\circ\text{C}$ and stirred overnight at room temperature, the reaction mixture was filtered to remove KCl. The resulting filtrate was then combined with the additional $\text{In}(\text{CH}_2\text{CMe}_3)_3$. Colorless crystals of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ formed after 7d. The product $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ (1.374 g, 1.708 mmol, 64% yield based on $\text{ClP}(\text{t-Bu})_2$) was finally isolated by filtration 72 d after the addition of $\text{In}(\text{CH}_2\text{CMe}_3)_3$. Characterization data were identical to that previously described.

Synthesis of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ by Thermolysis. The reagents, $\text{HP}(\text{t-Bu})_2$ (0.302 g, 2.06 mmol) and $\text{In}(\text{CH}_2\text{CMe}_3)_3$ (0.678 g, 2.07 mmol) were combined in a break-seal tube. After the tube was sealed, the mixture was heated at $120 - 160^\circ\text{C}$ for 2 d. The condensable gas was removed, measured and identified as 0.148 g of CMe_4 (2.06 mmol, 99% yield based on $\text{HP}(\text{t-Bu})_2$). The remaining crystalline solid was purified by sublimation at $110 - 125^\circ\text{C}$ ($\sim 10^{-2}$ mm Hg) and identified as $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ (0.698 g, 0.87 mmol, 84% yield based on $\text{HP}(\text{t-Bu})_2$). Characterization data were identical to that previously described.

Preparation of $\text{HP}(\text{t-Bu})_2$. A pentane solution of $\text{ClP}(\text{t-Bu})_2$ (1.034 g, 5.722 mmol) was added slowly to a mixture of KH (0.329 g, 8.21 mmol) and $\text{K}[\text{In}(\text{CH}_2\text{CMe}_3)_3\text{H}]$ (2.663 g, 7.229 mmol) in 10 mL pentane at $-40 \sim -50^\circ\text{C}$. After the mixture was warmed to room temperature and stirred for 10 h, pentane was removed by vacuum distillation at low temperature (0 to -20°C). The crude product was isolated by vacuum distillation into a -196°C trap. Final purification of $\text{HP}(\text{t-Bu})_2$ required removing a trace of pentane by vacuum distillation to yield $\text{HP}(\text{t-Bu})_2$ (0.492 g, 3.36 mmol, 59% yield based upon $\text{ClP}(\text{t-Bu})_2$). $\text{HP}(\text{t-Bu})_2$. ^1H NMR(δ , C_6D_6) 1.16 (d, $^3\text{J} = 12$ Hz), 3.14 (d, $^1\text{J} = 199$ Hz). $^{31}\text{P}\{^1\text{H}\}$ NMR (C_6D_6) δ 19.9 (s). ^{31}P NMR (C_6D_6) δ 19.9 (dm, $^1\text{J} = 199$ Hz, $^3\text{J} = 11$ Hz).

Freezing Point Depression Study of $(\text{Me}_3\text{CCH}_2)_3\text{In}\cdot\text{P}(\text{H})(\text{t-Bu})_2$ in Benzene. A tube was charged with $\text{In}(\text{CH}_2\text{CMe}_3)_3$ (0.0890 g, 0.271 mmol) and $\text{HP}(\text{t-Bu})_2$ (0.0374 g, 0.256 mmol) and then benzene (4.2995 g) was added. The freezing point of the resulting solution was measured 3 times. The solution was then diluted with 1.6433 g C_6H_6 . The average freezing points were used to calculate an observed molality and K_d : 0.0821 (0.011), 0.0611 (0.010). The experiment was repeated with $\text{In}(\text{CH}_2\text{CMe}_3)_3$ (0.0921 g, 0.281 mmol), $\text{HP}(\text{t-Bu})_2$ (0.0326 g, 0.223 mmol) and benzene (4.3892 g and 1.6435 g for dilution). Observed molality (K_d): 0.0798 (0.013), 0.0598 (0.013).

Reactions of $\text{In}(\text{CH}_2\text{CMe}_3)_3$ with $\text{HP}(\text{t-Bu})_2$ in Different Ratios. (a) A tube was charged with $\text{HP}(\text{t-Bu})_2$ (0.268 g, 1.83 mmol) and pentane solvent (1.968 g). The solution was then added to $\text{In}(\text{CH}_2\text{CMe}_3)_3$ (3.006 g, 9.159 mmol) at room temperature. The resulting solution was permitted to stand at ambient temperature. Colorless crystals were

observed after 6 d and isolated after 7 d. The product was isolated by removing the volatile compounds at 60 °C. The nonvolatile solid was identified as $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ (0.145 g, 0.181 mmol, 20% yield based on $\text{HP}(\text{t-Bu})_2$). (b) A second experiment which used similar concentrations of reactants $\text{HP}(\text{t-Bu})_2$ (0.306 g, 2.09 mmol), $\text{In}(\text{CH}_2\text{CMe}_3)_3$ (0.693 g, 2.11 mmol) in pentane (2.137 g) had the same experimental conditions as experiment a. No crystals were observed after 7 d. After all the volatile compounds were removed, only a faint trace of solid (less than 1 mg) remained in the flask. (c) The third experiment had $\text{In}(\text{CH}_2\text{CMe}_3)_3$ (0.702 g, 2.14 mmol), excess $\text{HP}(\text{t-Bu})_2$ (1.557 g, 10.65 mmol) and pentane (2.128 g). Again, no solid was observed after 7 d. After removing all the volatile compounds, no $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ remained in the flask.

Thermolysis of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$. Crystals of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ (0.618 g, 0.77 mmol) were sealed in a break-seal tube and heated at 245 °C for 1 h. The colorless volatile products (0.233 g) were then removed from the opened tube by vacuum distillation and identified as only hydrocarbons as no ^{31}P NMR lines were detected. The number and variety of ^1H and ^{13}C NMR lines in the spectrum of the volatile products suggested a mixture of unknown hydrocarbons. The material remaining in the tube was washed with 20 mL of pentane which removed an unknown brown, nonvolatile liquid. The resulting black powder was identified as InP (0.170 g, 1.17 mmol, 76% yield based on $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ used). **InP.** Anal. Found: C, 0.41; H, 0.04. X-ray Powder Diffraction [obsd (lit.²²): 3.56 (3.39), 2.91 (2.93), 2.06 (2.07), 1.76 (1.77), 1.69 (1.69), 1.47 (1.47), 1.35 (1.35), 1.31 (1.31). XPS Binding Energies (Atomic Concentration_{rel}) (Sample was supported on tape which was also detected with C 1s, 285 (83.09); O 1s 532 (16.91)): In

$d_{5/2}$, 494.5 (21.39); In $d_{3/2}$, 452.1 (21.39); P 2p, 128.5 (11.57); C 1s, 285.0 (51.01); O 1s, 532 (13.18);, Cl 2p, 199.4 (2.85).

Collection of X-Ray Diffraction Data for $[(\text{Me}_3\text{CCH}_2)\text{InP}(\text{t-Bu})_2]_2$. A well-formed colorless transparent crystal of dimensions $0.34 \times 0.30 \times 0.27$ mm was selected for the X-ray diffraction study. The crystal was prepared for the study by sealing it into a 0.3 mm thin walled capillary under strict anaerobic conditions. It was then mounted and aligned on a Siemens R3m/V four-circle single-crystal diffractometer. Details of the data collection²³ are presented in Table III.

The crystal belongs to the orthorhombic system. Determination of the space group was rendered difficult due to pseudo-symmetry which gave rise to a pseudo-face centering condition. The structure was finally solved successfully in space group Pbcn (No. 60)²⁴ where the systematic absences are $0k\ell$ for $k = 2n+1$, $h0\ell$ for $\ell = 2n+1$ and $hk0$ for $h+k = 2n+1$. In addition there is a systematic weakness for all reflections with $h+k = 2n+1$. (Only 460 of 2732 such reflections have $I > 3\sigma(I)$. The mean value for I/σ for reflections with $h+k = 2n+1$ is 3.0 as compared to a value of 23.0 for all reflections.) The successful solution of the structure in space group Pbcn reveals that the systematic weakness of the reflections with $h+k = 2n+1$ is a result of the two heaviest atoms in the structure (indium atoms with $Z = 49$) lying on a crystallographic two-fold axis at $x = 0$, $z = 3/4$ (Wyckoff notation c). These two atoms provide contributions only to the intensities of reflections with $h+k = 2n$.²⁴

[It should be noted that we earlier had concluded erroneously that the crystals had a C-centered orthorhombic Bravais lattice. Attempts to solve the structure in space groups

Cmc2₁ (No. 36) and Cmc_m (No. 63) were unsuccessful, but the structure was apparently solved in space group Ama2 (No. 40). Although refinement of data converged with $R = 4.20\%$, and the resulting molecular geometry appeared reasonable, the model had impossibly short intermolecular C(neopentyl)···C(neopentyl) contacts at 2.185\AA . It was this feature that led us back to consider a space group with a primitive orthorhombic lattice.]

Data were collected for one octant of the reciprocal sphere with $2\theta = 4.0\text{--}55.0^\circ$. Data were corrected for Lorentz and polarization effects but not for absorption. We estimate $\exp(-\mu\Delta t)$ to be 0.966, so that absorption can safely be ignored.

Solution and Refinement of the Structure of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$. All crystallographic calculations were carried out with the use of the Siemens SHELXTL PLUS program set.²⁵ The analytical scattering factors for neutral atoms were corrected for both the real and imaginary component ($\Delta f'$ and $i\Delta f''$) of anomalous dispersion components.²⁶

The structure was solved by a combination of direct methods and difference-Fourier syntheses. Positional and anisotropic thermal parameters were refined for all non-hydrogen atoms. Hydrogen atoms were included in idealized positions with $d(\text{C-H}) = 0.96\text{\AA}$ and the appropriate staggered tetrahedral geometry.²⁷ The isotropic thermal parameter of each of these hydrogen atoms was fixed equal to that of the carbon atom to which it was bonded. The final difference-Fourier map contained features ranging from $-0.51 \rightarrow +0.61\text{e}\text{\AA}^{-3}$. Refinement of the ordered model converged with $R = 6.68\%$ ($wR = 6.24\%$) for 183 parameters refined against all 4920 unique reflections ($R = 2.76\%$ and $wR = 3.47\%$ for those 2525 reflections with $|F_o| > 6\sigma(|F_o|)$). Final atomic coordinates are collected in Table IV.

Acknowledgements. This work was supported in part by the Office of Naval Research. Purchase of the Siemens R3m/V diffractometer was made possible by Grant No. 89-13733 from the Chemical Instrumentation Program of the National Science Foundation.

Supplementary Material Available. Anisotropic thermal parameters and calculated positions for all hydrogen atoms (2 pages). Ordering information is given on any current masthead page.

References

1. Cowley, A. H.; Benac, B. L.; Ekerdt, J. G.; Jones, R. A.; Kidd, K. B.; Lee, J. Y.; Miller, J. E. *J. Am. Chem. Soc.* **1988**, *110*, 6248.
2. Andrews, D. A.; Davies, G. J.; Bradley, D. C.; Faktor, M. M.; Frigo, D. M.; White, E. A. D. *Semicond. Sci. Technol.* **1988**, *3*, 1053.
3. Miller, J. E.; Kidd, K. B.; Cowley, A. H.; Jones, R. A.; Ekerdt, J. G.; Gysling, H. J.; Wernberg, A. A.; Blanton, T. N. *Chem. Mater.* **1990**, *2*, 589.
4. Miller, J. E.; Ekerdt, J. G. *Chem. Mater.* **1992**, *4*, 7.
5. Miller, J. E.; Mardones, M. A.; Nail, J. W.; Cowley, A. H.; Jones, R. H.; Ekerdt, J. G. *Chem. Mater.* **1992**, *4*, 447.
6. Coates, G. E.; Graham, J. J. *Chem. Soc.* **1963**, 233.
7. Beachley, O. T., Jr.; Coates, G. E. *J. Chem. Soc.* **1965**, 3241.
8. Maury, F.; Constant, G. *Polyhedron* **1984**, *3*, 581.
9. Beachley, O. T., Jr.; Kopasz, J. P.; Zhang, H.; Hunter, W. E.; Atwood, J. L. *J. Organomet. Chem.* **1987**, *325*, 69.
10. Banks, M. A.; Beachley, O. T., Jr.; Buttrey, L. A.; Churchill, M. R.; Fettingner, J. C. *Organometallics* **1991**, *10*, 1901.
11. Aitchison, K. A.; Backer-Dirks, J. D. J.; Bradley, D. C.; Faktor, M. M.; Fiegio, D. M.; Hursthouse, M. B.; Hussain, B.; Short, R. L. *J. Organomet. Chem.* **1989**, *366*, 11.
12. Alcock, N. W.; Degnan, I. E.; Wallbridge, M. G. H.; Powell, H. R.; McPartlin, M.; Sheldrick, G. M. *J. Organomet. Chem.* **1989**, *361*, C33.
13. Arif, A. M.; Benac, B. L.; Cowley, A. H.; Jones, R. A.; Kidd, K. B.; Nunn, C. M. *New J. Chem.* **1988**, *12*, 553.

14. Cowley, A. H.; Jones, R. A.; Mardones, M. A.; Nunn, C. M. *Organometallics* **1991**, *10*, 1635.
15. Wells, R. L.; McPhail, A. T.; Self, M. F. *Organometallics* **1992**, *11*, 221.
16. Wells, R. L., McPhail, A. T.; Jones, L. J.; Self, M. F. *Polyhedron* **1993**, *12*, 141.
17. Beachley, O. T., Jr.; Chao, S.-H. L.; Churchill, M. R.; See, R. F. *Organometallics* **1992**, *11*, 1486.
18. Beachley, O. T., Jr.; Tessier-Youngs, C. *Inorg. Chem.* **1979**, *18*, 3188.
19. Beachley, O. T., Jr.; Victoriano, L. *Inorg. Chem.* **1986**, *25*, 1948.
20. Beachley, O. T., Jr.; Spiegel, E. F.; Kopasz, J. P.; Rogers, R. D. *Organometallics* **1989**, *8*, 1915.
21. Shriver, D. F.; Drezdson, M. A. "The Manipulation of Air Sensitive Compounds", New York, 1986, p 38.
22. Joint Committee Powder Diffraction Standards, "Powder Diffraction File Search Manual", International Centre for Diffraction Data, Swarthmore, PA, File No. 32-452.
23. Churchill, M. R.; Lashewycz, R. A.; Rotella, F. J., *Inorg. Chem.* **1977**, *16*, 265.
24. International Tables for X-Ray Crystallography, Volume 1, Kynoch Press, Birmingham, England, 1965, pp 149 and 407.
25. Siemens SHELXTL PLUS Manual; 2nd Edition (1990). Siemens Analytical Instruments, Madison, Wisconsin.
26. International Tables for X-Ray Crystallography, Volume 4, Kynoch Press, Birmingham, England, 1974; pp 99-101 and 149-150.
27. Churchill, M. R., *Inorg. Chem.* **1973**, *12*, 1213.

Table I**Bond Lengths and Bond Angles for $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$.**

In(1)-P(1)	2.712 (1)	In(1)-C(31)	2.206 (5)
In(1)-P(1A)	2.712 (1)	In(1)-C(31A)	2.206 (5)
In(2)-P(1)	2.690 (1)	In(2)-C(41)	2.219 (4)
In(2)-P(1A)	2.690 (1)	In(2)-C(41A)	2.219 (4)
P(1)-C(11)	1.895 (4)	P(1)-C(21)	1.891 (4)
C(11)-C(12)	1.529 (6)	C(11)-C(13)	1.536 (6)
C(11)-C(14)	1.534 (6)	C(21)-C(22)	1.540 (6)
C(21)-C(23)	1.526 (6)	C(21)-C(24)	1.541 (6)
C(31)-C(32)	1.539 (6)	C(32)-C(33)	1.455 (8)
C(32)-C(34)	1.449 (8)	C(32)-C(35)	1.457 (7)
C(41)-C(42)	1.552 (6)	C(42)-C(43)	1.538 (6)
C(42)-C(44)	1.529 (6)	C(42)-C(45)	1.530 (6)

P(1)-In(1)-C(31)	116.6(1)	P(1)-In(1)-P(1A)	83.0(1)
C(31)-In(1)-P(1A)	109.1(1)	P(1)-In(1)-C(31A)	109.1(1)
C(31)-In(1)-C(31A)	117.7(2)	P(1A)-In(1)-C(31A)	116.6(1)
P(1)-In(2)-C(41)	109.7(1)	P(1)-In(2)-P(1A)	83.8(1)
C(41)-In(2)-P(1A)	119.2(1)	P(1)-In(2)-C(41A)	119.2(1)
C(41)-In(2)-C(41A)	112.7(2)	P(1A)-In(2)-C(41A)	109.7(1)
In(1)-P(1)-In(2)	96.6(1)	In(1)-P(1)-C(11)	116.7(1)
In(2)-P(1)-C(11)	110.6(1)	In(1)-P(1)-C(21)	109.6(1)
In(2)-P(1)-C(21)	112.6(1)	C(11)-P(1)-C(21)	110.2(2)
P(1)-C(11)-C(12)	105.9(3)	P(1)-C(11)-C(13)	113.8(3)
C(12)-C(11)-C(13)	109.1(4)	P(1)-C(11)-C(14)	111.9(3)
C(12)-C(11)-C(14)	107.6(3)	C(13)-C(11)-C(14)	108.2(3)
P(1)-C(21)-C(22)	105.8(3)	P(1)-C(21)-C(23)	112.6(3)
C(22)-C(21)-C(23)	108.1(3)	P(1)-C(21)-C(24)	112.4(3)
C(22)-C(21)-C(24)	108.8(3)	C(23)-C(21)-C(24)	109.0(3)
In(1)-C(31)-C(32)	124.9(3)	C(31)-C(32)-C(33)	110.8(5)
C(31)-C(32)-C(34)	109.5(5)	C(33)-C(32)-C(34)	106.8(8)
C(31)-C(32)-C(35)	115.0(4)	C(33)-C(32)-C(35)	106.2(5)
C(34)-C(32)-C(35)	108.2(6)	In(2)-C(41)-C(42)	124.2(3)
C(41)-C(42)-C(43)	109.0(3)	C(41)-C(42)-C(44)	111.0(3)
C(43)-C(42)-C(44)	108.0(4)	C(41)-C(42)-C(45)	112.6(3)
C(43)-C(42)-C(45)	108.1(4)	C(44)-C(42)-C(45)	108.0(4)

Table II**Comparisons of Bond Distances and Angles**

	<u>$[(\text{Me}_2\text{InP}(\text{t-Bu})_2)_2]_2$</u> ¹¹	<u>$[\text{Et}_2\text{InP}(\text{t-Bu})_2]_2$</u> ¹²	<u>$[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$</u>
In-C, Å	2.177 (6) 2.191 (6) 2.184 (Average)	2.170 (13) 2.176 (12) 2.173 (Average)	2.296 (5) 2.219 (4) 2.213 (Average)
In-P, Å	2.637 (4) 2.656 (4) 2.646 (Average)	2.635 (2)	2.712(1) 2.690 (1) 2.701 (Average)
In...In, Å	3.897	3.867	4.033
C-In-C,°	108.6 (3)	-----	117.7 (2) 112.7 (2)
P-In-P,°	85.2 (3)	85.6 (1)	83.8 (1) 83.0 (1) 83.4 (Average)
In-P-In,°	94.8 (2)	94.4 (1)	96.6 (1)

Table III**Experimental Data for the X-Ray Diffraction Study of $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$.****Crystal Data**

Empirical Formula	$\text{C}_{36} \text{H}_{80} \text{In}_2 \text{P}_2$
Crystal Size (mm)	0.34 x 0.3 x 0.27
Crystal System	Orthorhombic
Space Group	Pbcn
Unit Cell Dimensions	$a = 11.742(3) \text{ \AA}$ $b = 20.194(6) \text{ \AA}$ $c = 17.909(4) \text{ \AA}$
Volume	$4246(2) \text{ \AA}^3$
Z	4
Formula weight	804.6
Density(calc.)	1.259 Mg/m^3
Absorption Coefficient	1.163 mm^{-1}
F(000)	1696

Table III (Cont.)Data Collection

Diffractometer Used	Siemens R3m/V
Radiation	MoK α ($\lambda = 0.71073 \text{ \AA}$)
Temperature (K)	298 .
Monochromator	Highly oriented graphite crystal
2 θ Range	4.0 to 55.0 $^{\circ}$
Scan Type	2 θ - θ
Scan Speed	Constant; 3.00 $^{\circ}$ /min. in ω
Scan Range (ω)	0.45 $^{\circ}$ plus K α -separation
Background Measurement	Stationary crystal and stationary counter at beginning and end of scan, each for 25.0% of total scan time
Standard Reflections	3 measured every 97 reflections
Index Ranges	$0 \leq h \leq 15$, $-26 \leq k \leq 0$ $0 \leq l \leq 23$
Reflections Collected	5466
Independent Reflections	4920
Observed Reflections	2525 ($F > 6.0\sigma(F)$)

Table III (Cont.)Solution and Refinement

System Used	Siemens SHELXTL PLUS (VMS)
Solution	Direct Methods
Refinement Method	Full-Matrix Least-Squares
Quantity Minimized	$\sum w(F_o - F_c)^2$
Extinction Correction	$\chi = 0.00043(5)$, where $F^* = F [1 + 0.002\chi F^2 / \sin(2\theta)]^{-1/4}$
Hydrogen Atoms	Riding model, fixed isotropic U
Weighting Scheme	$w^{-1} = \sigma^2(F) + 0.0014F^2$
Number of Parameters refined	183
Final R indices (all data)	R = 6.68 %, wR = 6.24 %
R Indices (6.0 σ data)	R = 2.76 %, wR = 3.47 %
Goodness-of-Fit	0.97
Largest and Mean Δ/σ	0.003, 0.000
Data-to-Parameter Ratio	26.9:1
Largest Difference Peak	0.61 eÅ ⁻³
Largest Difference Hole	-0.51 eÅ ⁻³

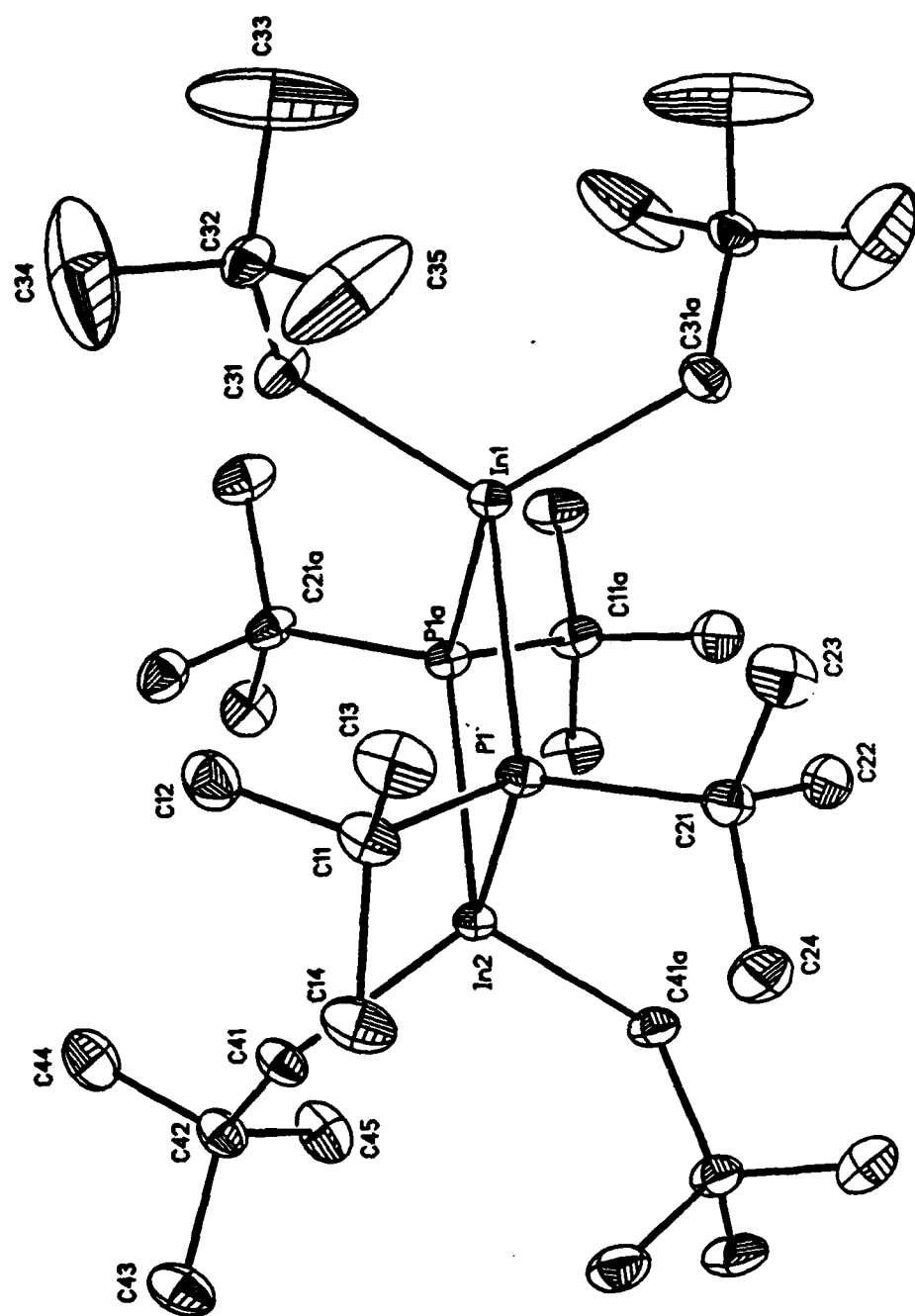
Table IV**Final Atomic Parameters for $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$.**Atomic coordinates ($\times 10^4$) and equivalent isotropic displacement coefficients ($\text{\AA}^2 \times 10^3$)

	x	y	z	U(eq)
In(1)	0	2741(1)	7500	33(1)
In(2)	0	744(1)	7500	33(1)
P(1)	1526(1)	1735(1)	7580(1)	34(1)
C(11)	2390(3)	1672(2)	8471(2)	47(1)
C(12)	1552(4)	1794(2)	9110(2)	60(2)
C(13)	3365(4)	2178(2)	8521(3)	67(2)
C(14)	2895(4)	978(2)	8577(3)	61(2)
C(21)	2503(3)	1764(2)	6742(2)	46(1)
C(22)	1728(4)	1682(2)	6055(2)	57(2)
C(23)	3123(3)	2426(2)	6670(3)	66(2)
C(24)	3387(3)	1200(2)	6746(3)	62(2)
C(31)	-297(3)	3306(2)	8536(3)	52(1)
C(32)	464(4)	3873(2)	8821(2)	46(1)
C(33)	-26(6)	4513(3)	8633(9)	231(8)
C(34)	535(9)	3843(5)	9628(4)	237(7)
C(35)	1619(5)	3876(3)	8523(5)	175(5)
C(41)	89(3)	135(2)	8530(2)	47(1)
C(42)	-770(4)	-422(2)	8729(2)	50(1)
C(43)	-108(4)	-1027(2)	9020(3)	73(2)
C(44)	-1586(4)	-197(2)	9344(3)	71(2)
C(45)	-1481(4)	-642(2)	8058(3)	69(2)

* Equivalent isotropic U defined as one third of the trace of the orthogonalized U_{ij} tensor

Caption to Figure

Figure 1. Labelling of atoms in the $[(\text{Me}_3\text{CCH}_2)_2\text{InP}(\text{t-Bu})_2]_2$ molecule. [ORTEP II diagram, with all hydrogen atoms omitted].



TECHNICAL REPORT DISTRIBUTION LIST - GENERAL

Office of Naval Research (2)*
Chemistry Division, Code 1113
800 North Quincy Street
Arlington, Virginia 22217-5000

Dr. James S. Murday (1)
Chemistry Division, Code 6100
Naval Research Laboratory
Washington, D.C. 20375-5000

Dr. Robert Green, Director (1)
Chemistry Division, Code 385
Naval Air Weapons Center
Weapons Division
China Lake, CA 93555-6001

Dr. Elek Lindner (1)
Naval Command, Control and Ocean
Surveillance Center
RDT&E Division
San Diego, CA 92152-5000

Dr. Bernard E. Douda (1)
Crane Division
Naval Surface Warfare Center
Crane, Indiana 47522-5000

Dr. Richard W. Drisko (1)
Naval Civil Engineering
Laboratory
Code L52
Port Hueneme, CA 93043

Dr. Harold H. Singerman (1)
Naval Surface Warfare Center
Carderock Division Detachment
Annapolis, MD 21402-1198

Dr. Eugene C. Fischer (1)
Code 2840
Naval Surface Warfare Center
Carderock Division Detachment
Annapolis, MD 21402-1198

Defense Technical Information
Center (2)
Building 5, Cameron Station
Alexandria, VA 22314

* Number of copies to forward